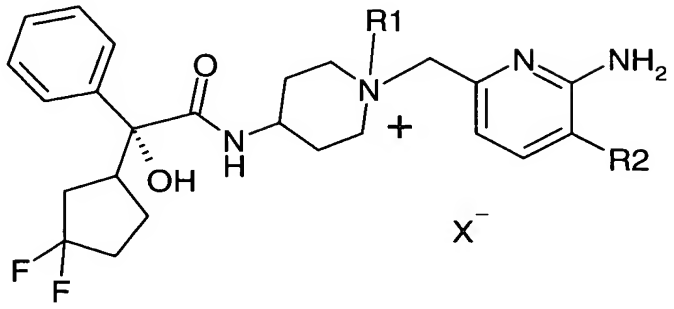
We claim:

1. A quaternary ammonium compound of formula I



and any stereoisomers thereof, wherein

R_1 is selected from C_1-C_6 alkyl, -CH_2-(C_1-C_4 alkenyl), and –CH_2-(C_1-C_6 alkynyl), each of which is optionally substituted with a group selected from phenyl, C_1-C_4 alkoxy, and hydroxyl;

R_2 is selected from H or OH; and

X represents an anion of a pharmaceutically acceptable acid.

2. The compound of claim 1, wherein X is selected from the group consisting of the anions of the following acids: tartaric, hydrochloric, hydrobromic, hydroiodic, sulfuric, phosphoric, nitric, citric, methanesulfonic, CH_3-$(CH_2)_n$-COOH where n is 0-4, HOOC-(CH_2)n-COOH where n is 1-4, HOOC-CH=CH-COOH, and benzoic.

3. The compound of claim 1, wherein X is selected from the group consisting of iodide, bromide, and chloride.

4. The compound of claim 1, wherein X is iodide.

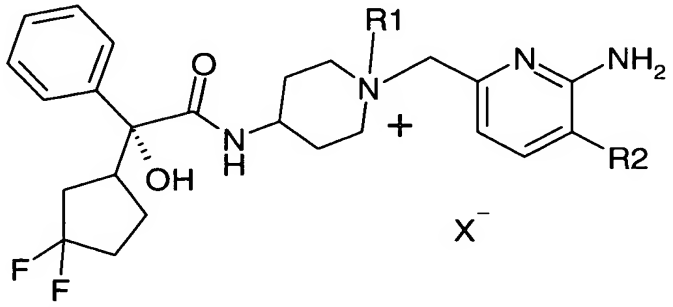
5. The compound of claim 1, wherein X is bromide.

6. The compound of claim 1, wherein X is chloride.

7. The compound of claim 1, wherein R_1 is methyl.

8. A compound (2R)-N-[1-(6-aminopyridin-2-ylmethyl)1-methylpiperdin-4-yl]-2-[(1R)-3,3,-difluorocyclopentyl]-2-hydroxy-2-phenylacetamide iodide.

9. A pharmaceutical composition comprising a pharmaceutical carrier and a therapeutically effective amount of a quaternary ammonium compound of formula I



and any stereoisomers thereof, wherein

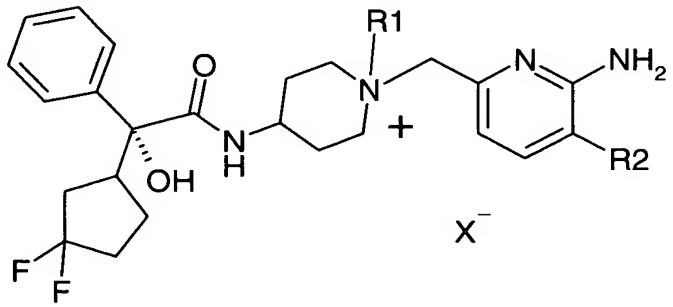
R_1 is selected from C_1-C_6 alkyl, -CH_2-(C_1-C_4 alkenyl), and –CH_2-(C_1-C_6 alkynyl), each of which is optionally substituted with a group selected from phenyl, C_1-C_4 alkoxy, and hydroxyl;

R_2 is selected from H or OH; and

X represents an anion of a pharmaceutically acceptable acid.

10. The method of treating a mammal for asthma, Chronic Obstructive Pulmonary Disease, allergic rhinitis, and infectious rhinitis, comprising:

administering a therapeutically effective amount of a quaternary ammonium compound of formula I, having the structure



and any stereoisomers thereof, wherein

R_1 is selected from C_1-C_6 alkyl, -CH_2-(C_1-C_4 alkenyl), and –CH_2-(C_1-C_6 alkynyl), each of which is optionally substituted with a group selected from phenyl,

C_1-C_4 alkoxy, and hydroxyl;

R_2 is selected from H or OH; and

X represents an anion of a pharmaceutically acceptable acid.